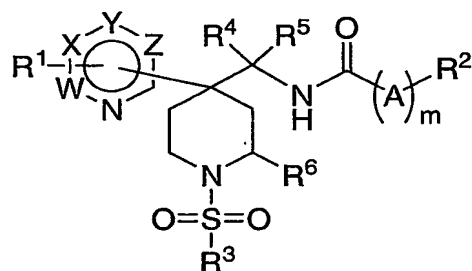


## WHAT IS CLAIMED IS:

1. A compound of the formula I:



I

wherein:

R<sup>1</sup> is selected from one or more of the groups consisting of:

- (1) hydrogen,
- (2) C<sub>1</sub>-6alkyl, which is unsubstituted or substituted with 1-6 halogen, hydroxy or phenyl,
- (3) -O-C<sub>1</sub>-6alkyl,
- (4) halogen,
- (5) phenyl, which is substituted with R<sup>2a</sup>, R<sup>2b</sup> and R<sup>2c</sup>,
- (6) heterocycle, which is substituted with R<sup>2a</sup>, R<sup>2b</sup> and R<sup>2c</sup>,
- (7) -CN,
- (8) -CO<sub>2</sub>R<sup>9</sup>,

wherein R<sup>9</sup> is independently selected from:

- (a) hydrogen,
- (b) -C<sub>1</sub>-6alkyl, which is unsubstituted or substituted with 1-6 fluoro,
- (c) benzyl, and
- (d) phenyl,
- (9) -SO<sub>2</sub>R<sup>9</sup>,
- (10) -SO<sub>2</sub>-NR<sup>10</sup>R<sup>11</sup>,

wherein R<sup>10</sup> and R<sup>11</sup> are independently selected from:

- (a) hydrogen,
- (b) -C<sub>1</sub>-6alkyl, which is unsubstituted or substituted with hydroxy, 1-6 fluoro or -NR<sup>12</sup>R<sup>13</sup>, where R<sup>12</sup> and R<sup>13</sup> are independently selected from hydrogen and -C<sub>1</sub>-6alkyl, and where R<sup>10</sup> and R<sup>11</sup> may be joined to form an azetidinyl ring,

- (c) -C<sub>3-6</sub>cycloalkyl, which is unsubstituted or substituted with hydroxy, 1-6 fluoro or -NR<sup>12</sup>R<sup>13</sup>,
- (d) benzyl,
- (e) phenyl, and

5 (11) -CONR<sup>10</sup>R<sup>11</sup>;

R<sup>2</sup> is selected from the group consisting of:

- (1) phenyl, which is substituted with R<sup>2a</sup>, R<sup>2b</sup> and R<sup>2c</sup>,
- (2) heterocycle, which is substituted with R<sup>2a</sup>, R<sup>2b</sup> and R<sup>2c</sup>,
- (3) C<sub>1-8</sub>alkyl, which is unsubstituted or substituted with 1-6 halogen, hydroxy,  
10 -NR<sup>10</sup>R<sup>11</sup>, phenyl or heterocycle, where the phenyl or heterocycle is substituted with R<sup>2a</sup>, R<sup>2b</sup> and R<sup>2c</sup>,
- (4) C<sub>3-6</sub>cycloalkyl, which is unsubstituted or substituted with 1-6 halogen, hydroxy or -NR<sup>10</sup>R<sup>11</sup>, and
- (5) -C<sub>1-6</sub>alkyl-(C<sub>3-6</sub>cycloalkyl), which is unsubstituted or substituted with 1-6 halogen,  
15 hydroxy or -NR<sup>10</sup>R<sup>11</sup>;

R<sup>2a</sup>, R<sup>2b</sup> and R<sup>2c</sup> are independently selected from the group consisting of:

- (1) hydrogen,
- (2) halogen,
- (3) -C<sub>1-6</sub>alkyl, which is unsubstituted or substituted with:
  - 20 (a) 1-6 halogen,
  - (b) phenyl,
  - (c) C<sub>3-6</sub>cycloalkyl, or
  - (d) -NR<sup>10</sup>R<sup>11</sup>,
- (4) -O-C<sub>1-6</sub>alkyl, which is unsubstituted or substituted with 1-6 halogen,
- 25 (5) hydroxy,
- (6) -SCF<sub>3</sub>,
- (7) -SCHF<sub>2</sub>,
- (8) -SCH<sub>3</sub>,
- (9) -CO<sub>2</sub>R<sup>9</sup>,
- 30 (10) -CN,
- (11) -SO<sub>2</sub>R<sup>9</sup>,
- (12) -SO<sub>2</sub>-NR<sup>10</sup>R<sup>11</sup>,
- (13) -NR<sup>10</sup>R<sup>11</sup>,
- (14) -CONR<sup>10</sup>R<sup>11</sup>, and

(15)  $-\text{NO}_2$ ;

$\text{R}^3$  is selected from the group consisting of:

- (1)  $\text{C}_{1-6}$ alkyl, which is unsubstituted or substituted with 1-6 halogen, hydroxyl,  $-\text{NR}^{10}\text{R}^{11}$ , or heterocycle, which is substituted with  $\text{R}^{2a}$ ,  $\text{R}^{2b}$  and  $\text{R}^{2c}$ ,
- 5 (2)  $\text{C}_{3-6}$ cycloalkyl, which is unsubstituted or substituted with 1-6 halogen, hydroxyl or  $-\text{NR}^{10}\text{R}^{11}$ ,
- (3)  $-\text{C}_{1-6}$ alkyl- $(\text{C}_{3-6}$ cycloalkyl), which is unsubstituted or substituted with 1-6 halogen, hydroxy or  $-\text{NR}^{10}\text{R}^{11}$ , and
- (4)  $-\text{NR}^{10}\text{R}^{11}$ , and
- 10 (5) heterocycle, which is substituted with  $\text{R}^{2a}$ ,  $\text{R}^{2b}$  and  $\text{R}^{2c}$ ;

$\text{R}^4$  and  $\text{R}^5$  are independently selected from the group consisting of:

- (1) hydrogen, and
- (2)  $\text{C}_{1-6}$ alkyl, which is unsubstituted or substituted with halogen or hydroxyl, or  $\text{R}^4$  and  $\text{R}^5$  taken together form a  $\text{C}_{3-6}$ cycloalkyl ring;

15  $\text{R}^6$  is selected from the group consisting of:

- (1) hydrogen, and
- (2)  $\text{C}_{1-6}$ alkyl;

W, X, Y and Z are independently selected from C or N, with the proviso that at least two of W, X, Y and Z are C, to form a pyridine, oxo-dihydropyridine, pyridazine, pyrimidine, pyrazine, 1,2,4-triazine  
20 or 1,3,5-triazine ring;

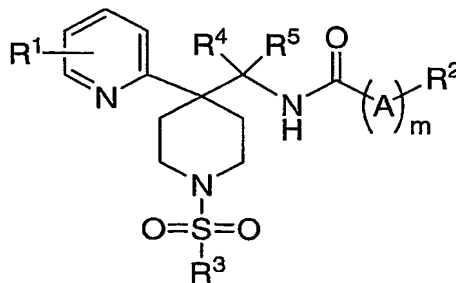
A is selected from the group consisting of:

- (1)  $-\text{O}-$ , and
- (2)  $-\text{NR}^{10}-$ ;

m is zero or one, whereby when m is zero  $\text{R}^2$  is attached directly to the carbonyl;

25 and pharmaceutically acceptable salts thereof.

2. The compound of Claim 1 of the formula Ia:



Ia

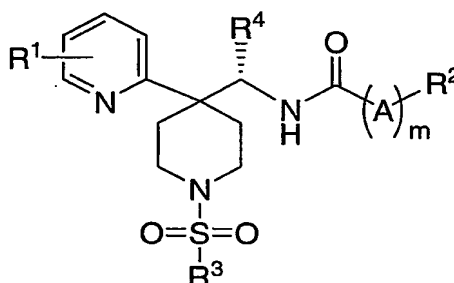
or a pharmaceutically acceptable salt thereof.

3. The compound of Claim 1 wherein  $R^1$  is selected from the group consisting of:

- (1) hydrogen,
- (2)  $C_{1-3}$ alkyl,
- (3) fluoro,
- (4)  $-CF_3$ ,
- (5)  $-morpholinyl$ , and
- (6)  $-O-C_{1-3}alkyl$ .

4. The compound of Claim 3 wherein  $R^1$  is hydrogen or methyl.

5. The compound of Claim 1 of the formula Ib:



Ib

wherein  $R^4$  is  $C_{1-6}$ alkyl; or a pharmaceutically acceptable salt thereof or an individual enantiomer or diastereomer thereof.

6. The compound of Claim 1 wherein  $R^4$  is  $C_{1-3}$ alkyl and  $R^5$  is hydrogen or  $C_{1-3}$ alkyl.

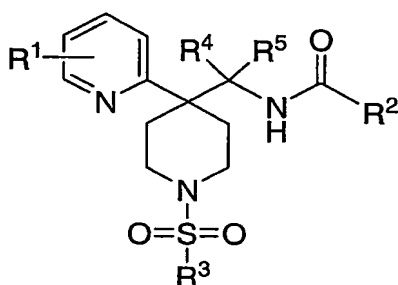
7. The compound of Claim 6 wherein  $R^4$  is  $C_{1-3}$ alkyl in the (S) configuration and  $R^5$  is hydrogen.

8. The compound of Claim 6 wherein  $R^4$  is methyl and  $R^5$  is hydrogen.

9. The compound of Claim 1 wherein  $R^4$  is methyl and  $R^5$  is methyl.

10. The compound of Claim 1 wherein  $R^4$  is hydrogen and  $R^5$  is hydrogen.

11. The compound of Claim 1 of the formula Ic:



Ic

or a pharmaceutically acceptable salt thereof.

12. The compound of Claim 1 wherein  $R^2$  is selected from the group consisting of:

- (1) phenyl, which is substituted with  $R^{2a}$ ,  $R^{2b}$  and  $R^{2c}$ ,
- (2) thienyl, which is substituted with  $R^{2a}$ ,  $R^{2b}$  and  $R^{2c}$ ,
- (3)  $C_1$ -alkyl, which is unsubstituted or substituted with 1-6 halogen, phenyl or  $-NR^{10}R^{11}$ , where the phenyl is substituted with  $R^{2a}$ ,  $R^{2b}$  and  $R^{2c}$ ,
- (4)  $C_3$ -cycloalkyl, which is unsubstituted or substituted with 1-6 halogen, hydroxy or  $-NR^{10}R^{11}$ , and

$R^{2a}$ ,  $R^{2b}$  and  $R^{2c}$  are independently selected from the group consisting of:

- (1) hydrogen,
- (2) halogen,
- (3)  $-C_1$ -alkyl,
- (4)  $-O-C_1$ -alkyl,
- (5)  $-CF_3$ ,
- (6)  $-OCF_3$ ,
- (7)  $-OCHF_2$ ,
- (8)  $-SCF_3$ ,
- (9)  $-SCHF_2$ , and
- (10)  $-NH_2$ .

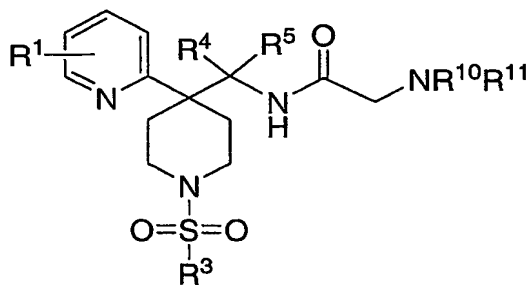
13. The compound of Claim 12 wherein  $R^2$  is phenyl or thienyl and  $R^{2a}$ ,  $R^{2b}$  and  $R^{2c}$  are independently selected from the group consisting of:

- (1) hydrogen,
- (2) halogen,
- (3)  $-C_{1-6}$ alkyl,
- (4)  $-O-C_{1-6}$ alkyl,
- (5)  $-CF_3$ ,
- (6)  $-OCF_3$ ,
- (7)  $-OCHF_2$ ,
- (8)  $-SCF_3$ ,
- (9)  $-SCHF_2$ , and
- (10)  $-NH_2$ .

14. The compound of Claim 13 wherein  $R^2$  is phenyl and  $R^{2a}$ ,  $R^{2b}$  and  $R^{2c}$  are independently selected from the group consisting of:

- (1) hydrogen,
- (2) fluoro,
- (3) chloro,
- (4) bromo,
- (5)  $-OCH_3$ ,
- (6)  $-CF_3$ , and
- (7)  $-NH_2$ .

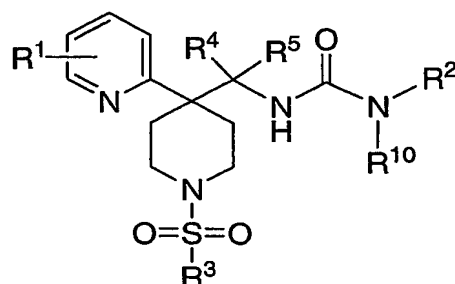
15. The compound of Claim 1 of the formula If:



If

or a pharmaceutically acceptable salt thereof.

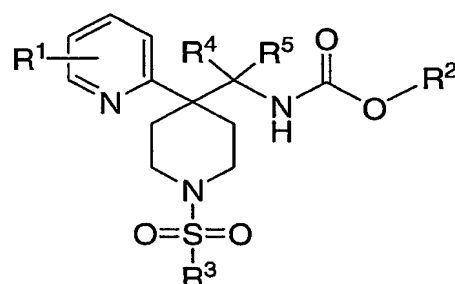
16. The compound of Claim 1 of the formula Ig:



Ig

5 or a pharmaceutically acceptable salt thereof.

17. The compound of Claim 1 of the formula Ih:



Ih

10 or a pharmaceutically acceptable salt thereof.

18. The compound of Claim 1 wherein R<sup>3</sup> is selected from C<sub>1</sub>-6alkyl, C<sub>1</sub>-6alkyl-fluoro, C<sub>3</sub>-6cycloalkyl, C<sub>1</sub>-6alkyl-cyclopropyl, -NH(C<sub>1</sub>-6alkyl), -N(C<sub>1</sub>-6alkyl)(C<sub>1</sub>-6alkyl) or azediny, which is unsubstituted or substituted with fluoro.

15

19. The compound of Claim 18 wherein R<sup>3</sup> is -CH<sub>2</sub>CH<sub>3</sub>.

20. The compound of Claim 18 wherein R<sup>3</sup> is -(CH<sub>2</sub>)<sub>2</sub>CH<sub>3</sub>.

20

21. The compound of Claim 18 wherein R<sup>3</sup> is -CH<sub>2</sub>-cyclopropyl.

22. A compound which is selected from the group consisting of:

- 2-chloro-3,6-difluoro-N-{{1-(propylsulfonyl)-4-pyridin-2-yl}piperidin-4-yl}methyl}benzamide;
- 2,4-dichloro-N-{{4-(6-methylpyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl}methyl}benzamide;
- 2,4-dichloro-N-{{(1S)-1-[1-(ethylsulfonyl)-4-pyridin-2-yl}piperidin-4-yl}ethyl}benzamide;
- 5 2-chloro-3,6-difluoro-N-{{1-methyl-1-[1-(propylsulfonyl)-4-pyridin-2-yl}piperidin-4-yl}ethyl}benzamide;
- 2-chloro-N-{{1-(propylsulfonyl)-4-pyridin-2-yl}piperidin-4-yl}methyl}benzamide;
- 2,6-dichloro-N-{{1-(propylsulfonyl)-4-pyridin-2-yl}piperidin-4-yl}methyl}-benzamide;
- 2-bromo-N-{{1-(propylsulfonyl)-4-pyridin-2-yl}piperidin-4-yl}methyl}benzamide;
- 2,4-dichloro-N-{{1-(propylsulfonyl)-4-pyridin-2-yl}piperidin-4-yl}methyl}benzamide;
- 10 2-chloro-6-fluoro-N-{{1-(propylsulfonyl)-4-pyridin-2-yl}piperidin-4-yl}methyl}benzamide;
- 2-amino-6-chloro-N-{{1-(propylsulfonyl)-4-pyridin-2-yl}piperidin-4-yl}methyl}benzamide;
- 2-fluoro-6-methoxy-N-{{1-(propylsulfonyl)-4-pyridin-2-yl}piperidin-4-yl}methyl}benzamide;
- 2-chloro-N-{{4-(6-methylpyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl}methyl}benzamide;
- 2-fluoro-N-{{4-(6-methylpyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl}methyl}-6-
- 15 (trifluoromethyl)benzamide;
- 2,6-difluoro-N-{{4-(6-methylpyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl}methyl}benzamide;
- 2-chloro-6-fluoro-N-{{4-(6-methylpyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl}methyl}benzamide;
- 2,6-dichloro-N-{{4-(6-methylpyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl}methyl}benzamide;
- 2-chloro-3,6-difluoro-N-{{4-(6-methylpyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl}methyl}benzamide;
- 20 2-chloro-4-fluoro-N-{{4-(6-methylpyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl}methyl}benzamide;
- 4-chloro-2-fluoro-N-{{4-(6-methylpyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl}methyl}benzamide;
- 2,4-dichloro-N-{{4-(4-methylpyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl}methyl}benzamide
- 2,4-dichloro-N-{{1-(methylsulfonyl)-4-pyridin-2-yl}piperidin-4-yl}methyl}benzamide;
- 2,4-dichloro-N-{{1-(isopropylsulfonyl)-4-pyridin-2-yl}piperidin-4-yl}methyl}benzamide
- 25 2,4-dichloro-N-{{1-(ethylsulfonyl)-4-pyridin-2-yl}piperidin-4-yl}methyl}benzamide;
- 2,4-dichloro-N-{{1-(cyclopropylsulfonyl)-4-pyridin-2-yl}piperidin-4-yl}methyl}benzamide;
- 2,4-dichloro-N-{{1-(propylsulfonyl)-4-pyridin-3-yl}piperidin-4-yl}methyl}benzamide;
- 2,6-dichloro-N-{{1-(propylsulfonyl)-4-pyridin-3-yl}piperidin-4-yl}methyl}benzamide
- 2,4-dichloro-N-{{1-(propylsulfonyl)-4-pyridin-4-yl}piperidin-4-yl}methyl}benzamide;
- 30 2-chloro-6-fluoro-N-{{1-(propylsulfonyl)-4-pyridin-4-yl}piperidin-4-yl}methyl}benzamide;
- 2,4-dichloro-N-{{1-[(dimethylamino)sulfonyl]-4-pyridin-2-yl}piperidin-4-yl}methyl}benzamide;
- 4,4,4-trifluoro-3-methyl-N-{{4-(6-methylpyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl}methyl}butanamide;



- 2-chloro-6-fluoro-N-{[4-(6-morpholin-4-ylpyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl]methyl}benzamide;  
2,4-dichloro-N-{[4-(6-morpholin-4-ylpyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl]methyl}benzamide;  
2,4,5-trifluoro-N-{[4-(6-methoxypyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl]methyl}benzamide;  
5 2,4-dichloro-5-fluoro-N-{[4-(6-methoxypyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl]methyl}benzamide;  
N-{[4-(6-methylpyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl]methyl}cyclohexanecarboxamide;  
2-chloro-N-{[1-(cyclopropylsulfonyl)-4-(6-methylpyridin-2-yl)piperidin-4-yl]methyl}-3,6-difluorobenzamide;  
10 N-{[1-(ethylsulfonyl)-4-pyridin-2-ylpiperidin-4-yl]methyl}-2,4-difluorobenzamide;  
N-(sec-butyl)-N'-{[1-(propylsulfonyl)-4-pyridin-2-ylpiperidin-4-yl]methyl}urea;  
N-(4-bromophenyl)-N'-{[1-(propylsulfonyl)-4-pyridin-2-ylpiperidin-4-yl]methyl}urea;  
3-fluorobenzyl {[1-(propylsulfonyl)-4-pyridin-2-ylpiperidin-4-yl]methyl}carbamate;  
2-chlorobenzyl {[1-(propylsulfonyl)-4-pyridin-2-ylpiperidin-4-yl]methyl}carbamate;  
15 2,4-dichloro-N-{1-methyl-1-[1-(propylsulfonyl)-4-pyridin-2-ylpiperidin-4-yl]ethyl}benzamide;  
2-chloro-3,6-difluoro-N-{1-methyl-1-[1-(propylsulfonyl)-4-pyridin-2-ylpiperidin-4-yl]ethyl}benzamide;  
N-{1-methyl-1-[1-(propylsulfonyl)-4-pyridin-2-ylpiperidin-4-yl]ethyl}-2-(trifluoromethoxy)benzamide;  
2,4-dichloro-N-{1-methyl-1-[4-(6-methylpyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl]ethyl}benzamide;  
2-chloro-3,6-difluoro-N-{1-methyl-1-[4-(6-methylpyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl]ethyl}benzamide;  
20 2,4-dichloro-N-{(1S)-1-[1-(propylsulfonyl)-4-pyridin-2-ylpiperidin-4-yl]ethyl}benzamide;  
2-chloro-3,6-difluoro-N-{(1S)-1-[1-(propylsulfonyl)-4-pyridin-2-ylpiperidin-4-yl]ethyl}benzamide;  
2,4-dichloro-N-{(1S)-1-[4-(6-methylpyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl]ethyl}benzamide;  
2-chloro-3,6-difluoro-N-{(1S)-1-[4-(6-methylpyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl]ethyl}benzamide;  
25 4,4,4-trifluoro-3-methyl-N-{(1S)-1-[1-(propylsulfonyl)-4-pyridin-2-ylpiperidin-4-yl]ethyl}butanamide;  
N-{(1S)-1-[1-(propylsulfonyl)-4-pyridin-2-ylpiperidin-4-yl]ethyl}thiophene-3-carboxamide;  
N-{(1S)-1-[1-(propylsulfonyl)-4-pyridin-2-ylpiperidin-4-yl]ethyl}cyclopentane-carboxamide;  
N-{(1S)-1-[1-(propylsulfonyl)-4-pyridin-2-ylpiperidin-4-yl]ethyl}cyclohexane-carboxamide;  
30 2-ethyl-N-{(1S)-1-[1-(propylsulfonyl)-4-pyridin-2-ylpiperidin-4-yl]ethyl}butanamide;  
2-methyl-N-{(1S)-1-[1-(propylsulfonyl)-4-pyridin-2-ylpiperidin-4-yl]ethyl}butanamide;  
3,3,3-trifluoro-N-{(1S)-1-[4-(6-methylpyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl]ethyl}propanamide;  
2,5-dichloro-N-{(1S)-1-[4-(6-methylpyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl]ethyl}thiophene-3-carboxamide;

- 4-bromo-N-((1S)-1-[4-(6-methylpyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl]ethyl)thiophene-3-carboxamide;
- 2,5-dichloro-N-((1S)-1-[1-(propylsulfonyl)-4-pyridin-2-ylpiperidin-4-yl]ethyl)thiophene-3-carboxamide;
- 4-bromo-N-((1S)-1-[1-(propylsulfonyl)-4-pyridin-2-ylpiperidin-4-yl]ethyl)thiophene-3-carboxamide;
- 5 2-chloro-N-([1-(propylsulfonyl)-4-pyrimidin-4-ylpiperidin-4-yl]methyl)benzamide;
- 2,4-dichloro-N-([1-(propylsulfonyl)-4-pyrimidin-4-ylpiperidin-4-yl]methyl)benzamide;
- 2-chloro-3,6-difluoro-N-([1-(propylsulfonyl)-4-pyrimidin-4-ylpiperidin-4-yl]methyl)benzamide;
- 2,4-dichloro-5-fluoro-N-([1-(propylsulfonyl)-4-pyrimidin-4-ylpiperidin-4-yl]methyl)benzamide;
- 2-chloro-N-((1S)-1-[1-(ethylsulfonyl)-4-pyridin-2-ylpiperidin-4-yl]ethyl)benzamide;
- 10 2-chloro-N-((1S)-1-[1-(ethylsulfonyl)-4-pyridin-2-ylpiperidin-4-yl]ethyl)-3,6-difluorobenzamide;
- 2-chloro-N-((1S)-1-[1-(ethylsulfonyl)-4-(6-methylpyridin-2-yl)piperidin-4-yl]ethyl)-3,6-difluorobenzamide;
- 2,4-dichloro-N-((1S)-1-[1-(ethylsulfonyl)-4-(6-methylpyridin-2-yl)piperidin-4-yl]ethyl)benzamide;
- 2,4-dichloro-N-([1-methyl-1-[1-(azetidinesulfonyl)-4-pyridin-2-ylpiperidin-4-yl]ethyl)benzamide;
- 15 2,4-dichloro-N-([1-methyl-1-[1-(3-fluoroazetidinesulfonyl)-4-pyridin-2-ylpiperidin-4-yl]ethyl)benzamide;
- 2,4-dichloro-N-([1-methyl-1-[1-(3-fluoroazetidinesulfonyl)-4-pyridin-2-ylpiperidin-4-yl]ethyl)benzamide;
- 2,4-dichloro-N-([1-methyl-1-[1-(azetidinesulfonyl)-4-(3-fluoropyridin-2-yl)piperidin-4-yl]ethyl)benzamide;
- 2,4-dichloro-N-([1-methyl-1-[1-(ethylaminosulfonyl)-4-pyridin-2-ylpiperidin-4-yl]ethyl)benzamide;
- 20 2,4-dichloro-N-([1-[(ethylamino)sulfonyl]-4-(3-fluoropyridin-2-yl)piperidin-4-yl]methyl)benzamide;
- 2,4-dichloro-N-([1-[1-(ethylaminosulfonyl)-4-pyridin-2-ylpiperidin-4-yl]ethyl)benzamide;
- 2,4-dichloro-N-([1-[1-(ethylaminosulfonyl)-4-(3-fluoropyridin-2-yl)piperidin-4-yl]ethyl)benzamide;
- 2,4-dichloro-N-([4-(3-methylpyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl]methyl)benzamide;
- 2,4-dichloro-N-([1-(propylsulfonyl)-4-[6-(trifluoromethyl)pyridin-2-yl]piperidin-4-yl]methyl)benzamide;
- 25 2,4-dichloro-N-([1-[(cyclopropylmethyl)sulfonyl]-4-(3-methylpyridin-2-yl)piperidin-4-yl]methyl)-benzamide;
- 2,4-dichloro-N-([1-(propylsulfonyl)-4-[4-(trifluoromethyl)pyridin-2-yl]piperidin-4-yl]methyl)benzamide;
- 2,4-dichloro-N-([4-(3-chloropyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl]methyl)benzamide;
- 2,4-dichloro-N-([4-(3-methoxypyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl]methyl)benzamide;
- 30 2-chloro-N-([4-(3-chloropyridin-2-yl)-1-(ethylsulfonyl)piperidin-4-yl]methyl)-3,6-difluorobenzamide;
- 2,4-dichloro-N-([1-[(cyclopropylmethyl)sulfonyl]-4-pyridin-2-ylpiperidin-4-yl]methyl)benzamide;
- methyl [(4-[(2,4-dichlorobenzoyl)amino]methyl)-4-pyridin-2-ylpiperidin-1-yl)sulfonyl]acetate;
- 2,4-dichloro-N-([1-(ethylsulfonyl)-4-pyridin-2-ylpiperidin-4-yl]methyl)benzamide;
- 2,4-dichloro-N-([1-(propylsulfonyl)-4-pyrazin-2-ylpiperidin-4-yl]methyl)benzamide;

- 2,4-dichloro-N-(1-{1-[(3-fluoropropyl)sulfonyl]-4-pyridin-2-yl}piperidin-4-yl)ethyl)benzamide;  
 2,4-dichloro-N-({1-[(3-fluoropropyl)sulfonyl]-4-pyridin-2-yl}piperidin-4-yl)methyl)benzamide;  
 2,4-dichloro-N-({1-[(3-fluoropropyl)sulfonyl]-4-(3-fluoropyridin-2-yl)}piperidin-4-yl)methyl)benzamide;  
 2,4-dichloro-N-({1-[(cyclopropylmethyl)sulfonyl]-4-(3-fluoropyridin-2-yl)}piperidin-4-yl)methyl)  
 5 benzamide;  
 2,4-dichloro-N-({4-(3-fluoropyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl)methyl}benzamide;  
 2-chloro-3,5-difluoro-N-({1-[(cyclopropylmethyl)sulfonyl]-4-(3-fluoropyridin-2-yl)}piperidin-4-  
 yl)methyl}benzamide;  
 2,4-dichloro-5-fluoro-N-({1-[(cyclopropylmethyl)sulfonyl]-4-(3-fluoropyridin-2-yl)}piperidin-4-  
 10 yl)methyl}benzamide;  
 2,4-dichloro-N-({1-[4-(3-fluoropyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl]ethyl}benzamide;  
 2,4-dichloro-N-({4-(6-trifluoromethylpyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl)methyl}benzamide;  
 4-bromo-2-chloro-N-({1-(ethylsulfonyl)-4-(3-fluoropyridin-2-yl)-piperidin-4-yl)methyl}benzamide;  
 N-(3-fluorobenzyl)-N'-({1-(propylsulfonyl)-4-(6-trifluoromethylpyridin-2-yl)}piperidin-4-yl)methyl}urea;  
 15 2,4-dichloro-N-({1-[1-(ethylsulfonyl)-4-(3-fluoropyridin-2-yl)-piperidin-4-yl]ethyl}benzamide;  
 2-bromo-4-fluoro-N-({1-(ethylsulfonyl)-4-(3-fluoropyridin-2-yl)-piperidin-4-yl)methyl}benzamide;  
 2-chloro-3,6-difluoro-N-({1-(ethylsulfonyl)-4-(3-fluoropyridin-2-yl)-piperidin-4-yl)methyl}benzamide;  
 2,4-dichloro-N-({1-(ethylsulfonyl)-4-(3-trifluoromethylpyridin-2-yl)-piperidin-4-yl)methyl}benzamide;  
 2,4-dichloro-N-({1-[(3-fluoropropyl)sulfonyl]-4-(3-trifluoromethylpyridin-2-yl}piperidin-4-  
 20 yl)methyl}benzamide;  
 2,4,6-trifluoro-N-({1-(ethylsulfonyl)-4-(3-trifluoromethylpyridin-2-yl)-piperidin-4-  
 yl)methyl}benzamide;  
 N-(sec-butyl)-N'-({1-(propylsulfonyl)-4-(6-trifluoromethylpyridin-2-yl)}piperidin-4-yl)methyl}urea;  
 2,4-dichloro-N-({1-(ethylsulfonyl)-4-(3-trifluoromethylpyridin-2-yl)-piperidin-4-yl)methyl}benzamide;  
 25 2,4-dichloro-N-({1-[4-(3-methylpyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl]ethyl}benzamide;  
 N-({4-(3-bromopyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl)methyl}-2,4-dichlorobenzamide;  
 2,4-dichloro-N-({4-(6-fluoropyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl)methyl}benzamide;  
 2,4-dichloro-N-({4-(6-chloropyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl)methyl}benzamide;  
 2,4-dichloro-N-({4-(6-oxo-1,6-dihydropyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl)methyl}benzamide;  
 30 2,4-dichloro-N-({4-(3-hydroxypyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl)methyl}benzamide;  
 2,4-dichloro-N-({(2[R,S],4[S,R])-2-methyl-1-(propylsulfonyl)-4-pyridin-2-yl}piperidin-4-  
 yl)methyl}benzamide;  
 2,4-dichloro-N-({1-(ethylsulfonyl)-4-[3-(trifluoromethyl)pyridin-2-yl]}piperidin-4-yl)methyl}benzamide;  
 or a pharmaceutically acceptable salt thereof.

23. A pharmaceutical composition which comprises a pharmaceutically acceptable carrier and a compound of Claim 1 or a pharmaceutically acceptable salt thereof.

5 24. A method for inhibiting the glycine transporter GlyT1 in a mammal in need thereof which comprises the administration of an effective amount of the compound of Claim 1 or a pharmaceutically acceptable salt thereof.

10 25. A method for the manufacture of a medicament for inhibiting the glycine transporter GlyT1 in a mammal in need thereof comprising combining the compound of Claim 1 or a pharmaceutically acceptable salt thereof with a pharmaceutical carrier or diluent.

15 26. A method for treating a neurological and psychiatric disorders associated with glycinergic or glutamatergic neurotransmission dysfunction in a mammalian patient in need thereof which comprises administering to the patient a therapeutically effective amount of a compound of Claim 1 or a pharmaceutically acceptable salt thereof.

20 27. A method for treating schizophrenia in a mammalian patient in need thereof which comprises administering to the patient a therapeutically effective amount of a compound of Claim 1 or a pharmaceutically acceptable salt thereof.